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DISTRIBUTION OF RADIOACTIVITY IN SEA WATER AND MARINE ORGANISMS FOLLOWING AN UNDERWATER NUCLEAR DETONATION AT THE ENIWETOK TEST SITE IN 1958

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ABSTRACT

An investigation of the radioactive contamination of the water and marine organisms in and near the Eniwetok Test Site was conducted during May 1958, shortly before and immediately following an underwater nuclear detonation. At the end of three and one-half days the boundaries of the radioactive water mass extended beyond the survey area, 50 miles to the west of Eniwetok Atoll, and to a depth of at least 300 meters. The plankton contained high levels of Np²³⁹, Mo⁹⁹-Tc^{99m}, Te¹³²-I¹³², and U²³⁷. Also present in lower amounts were Ce¹⁴¹-Pr¹⁴¹, Ru¹⁰³-Rh¹⁰³, Ba¹⁴⁰-La¹⁴⁰, Zr⁹⁵-Nb⁹⁵, and Ce¹⁴⁴-Pr¹⁴⁴. Ru¹⁰⁶-Rh¹⁰⁶ and Ru¹⁰⁵-Rh¹⁰⁵ were detected once in each of two samples.

Whole fish samples contained essentially the same radioisotopes as the plankton. Shrimp and squid contained high levels of Np^{239} and Te^{132} - I^{132} but no detectable Mo^{99} - Tc^{99m} .

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INTRODUCTION

Studies were made by the United States in 1955 and 1956

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and by Japan in 1954 and 1956 to determine the extent of the residual radioactive contamination in the western Pacific Ocean resulting from the nuclear tests at the Eniwetok Test Site. These studies were concerned with radioactive fallout material produced by the detonation of nuclear devices on land or in very shallow water.

The present investigation was conducted during the 1958 test series (Hardtack) and was designed primarily to determine the uptake of the short-lived isotopes by marine organisms and to outline the mass of radioactive water immediately following the underwater detonation of a nuclear device (Wahoo)

The survey was divided into two phases; the first, a predetonation survey, was designed to evaluate the radioactivity contributed by the earlier tests of the series; the second, a post detonation survey, covered the four-day period immediately following the Wahoo detonation.

The Hydrographic Office of the U.S. Navy assigned the USS

Rehoboth (AGS-50) to assist the Office of Naval Research in a research

program related to the Wahoo event. The Atomic Energy Commission supplied part of the funds for the operational cost of the Rehoboth and in return the Rehoboth was made available for use by the University of Washington Laboratory of Radiation Biology, an AEC contractor, for the period May 3, 1958 to May 20, 1958, to conduct a radiobiological survey.

Eight U. S. Navy Hydrographic Office personnel, under the direction of Mr. Alfred W. Anderson, made oceanographic surveys during both phases of the investigation and provided the authors with the water samples from four locations for measurement of radioactivity.

A radiation monitoring team from the Radiation Technical Division of the Naval Medical Research Institute, under the direction of Lt.

J. W. Duckworth, measured the radioactivity of water at the surface, at the intake to the ship's evaporators, and at designated depths from the surface down to 600 feet. The surface measurements were made with a survey meter and the measurements of the inflow water to the evaporators were made with a specially designed ion-chamber that was fitted around the intake line. For measurements at depth a gamma scintillation-type probe with a continuous recording device was used. The probe was a modified "well-logging" type provided by the AEC Health and Safety Laboratory, New York Operations Office. A report of these measurements is being prepared for publication by the Naval Medical Research Institute.

The measurement of radioactivity of the surface water (initial radiation survey) was conducted with the assistance of personnel from the Laboratory of Radiation Biology (University of Washington) and from the Radiation Safety Section of the Los Alamos Scientific Laboratory under the direction of Maj. Gordon Jacks.

The radiobiological survey extended from 10° 56' N to 11° 25' N and from 161° 16' E to 162° 10' E, the area southwest of Eniwetok Atoll (Fig. 1). The survey included thirty-five plankton stations and four water stations. Eight mid-water trawl collections also were made during the period. Details of these collections describing location, depth of sample, temperature, and depth of thermocline are given in Table 1 and Appendix Tables A, B, and C, and the locations of the stations are shown in Figures 1, 2, and 3. Details of the work aboard ship and the methods of collection were similar to those described by Donaldson et al. and Seymour et al.

MATERIALS AND METHODS

Plankton

Plankton samples were collected with a 1/2-meter plankton net of No. 6 mesh while the ship was drifting at a rate of approximately 0.5 to 1.8 knots per hour. The net, hauled by a 3/16-inch wire cable on a

Table 1. Gross beta activity and temperature of the water obtained with Nansen casts. Values expressed in d/m/l in the particulate matter and in the water (filterable portion less K⁴⁰ at time of collection.

					d/m/l**	
	Sample	Depth	Temp.	Particulat	е	
Station	no.	(meters)	°C	matter***	Water	Total
						_
H-7	106	0	27.82	120	7,400	7,520
Collected	107	8	(27.84)*	170	600	770
5/5/58	108	16	27.81	0	9,000	9,000
1408	109	24	27.81	0	0	0
11°21.8' N;	110	39	27.77	0	850	850
162°09.1'E	111	59	27.77	0	0	0
	112	79	26.69	0	0	0
	113	119	24.68	` 0	0	0
	114	160	17.85	0	0	0
	115	201	13.21	0	0	0
	116	244	11.07	0	0	0
	117	337	(9.70) *	0	0	0
	118	334	10.04	0	0	0
	119	416	(9.51)*	0	0	0
	120	501	8.17	0	0	0
	121	586	7.22	0	0	0
	122	675	6.27	0	0	0
	123	857	5.12	0	0	0
	124	1039	4.15	0	0	0
W-1	513	0	27.66	64,000	160,000	224,000
Collected	514	5	27.69	62,000	120,000	182,000
5/16/58	515	25	27.69	10,000	73,000	83,000
1910	516	50	27.58	22,000	30,000	52,000
11°19.3'N;	517	100	26.76	3,800	22,000	25,800
162°09.4'E	518	200	15.79	1,900	30,000	31,900
	519	300	10.71	3,100	(lost)	-

^{*} Doubtful.

^{**} Correction factor for self absorption based on K⁴⁰.

^{*** &}gt; 0.45 μ .

Table 1. - continued

d/m/l** Particulate Sample Depth Temp. matter*** Station no. (meters) °C Water Total W-2 520 0 27.63 30,000 165,000 195,000 24,000 152,000 176,000 29 27,58 Collected 521 19,000 141,000 160,000 48 27.60 5/17/58 522 69,900 1653 523 72 27.56 9,900 60,000 27.78 7,600 24,000 31,600 11°20.5'N 524 96 740 26,000 26,740 162°06.9'E 22.31 525 145 32,000 34,500 526 293 10.08 2,500 12,000 15,500 W-3 0 27.60 ` 3,500 527 Collected 528 30 27.55 2,300 14,000 16,300 3,000 11,000 14,000 5/18/58 50 27.56 529 75 27.49 3,400 11,000 14, 400 1348 530 65,000 74,000 139,000 11°20.9'N 531 100 27.34 13,000 18,300 21,98 5,300 162°10.6'E 532 150 49,000 533 300 09.88 29,000 78,000

^{*} Doubtful.

^{**} Correction factor for self absorption based on K⁴⁰.

^{*** &}gt; 0.45μ .

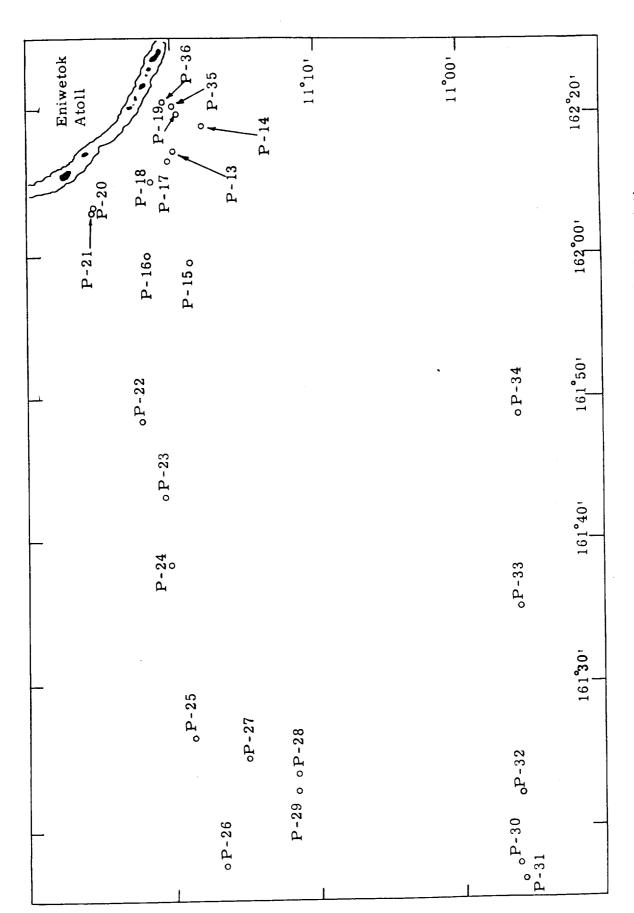
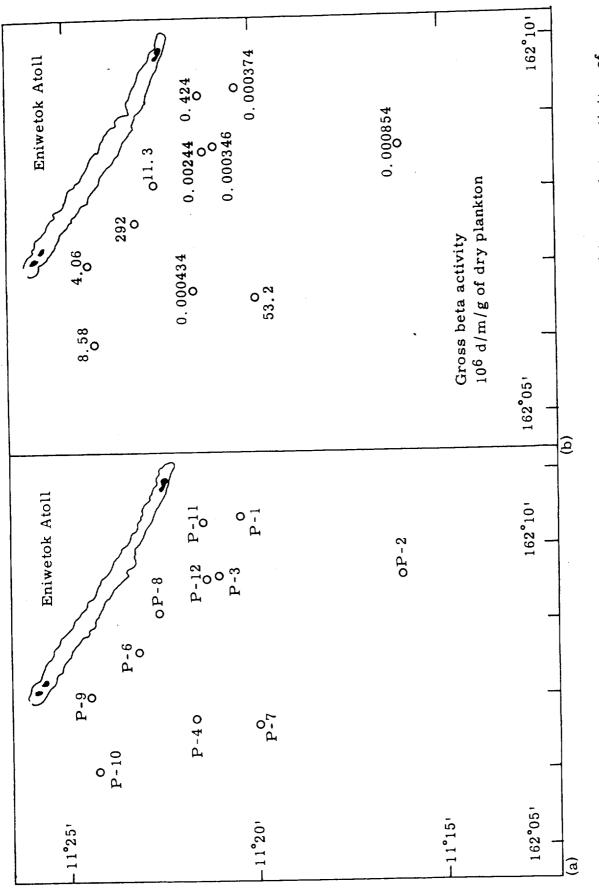
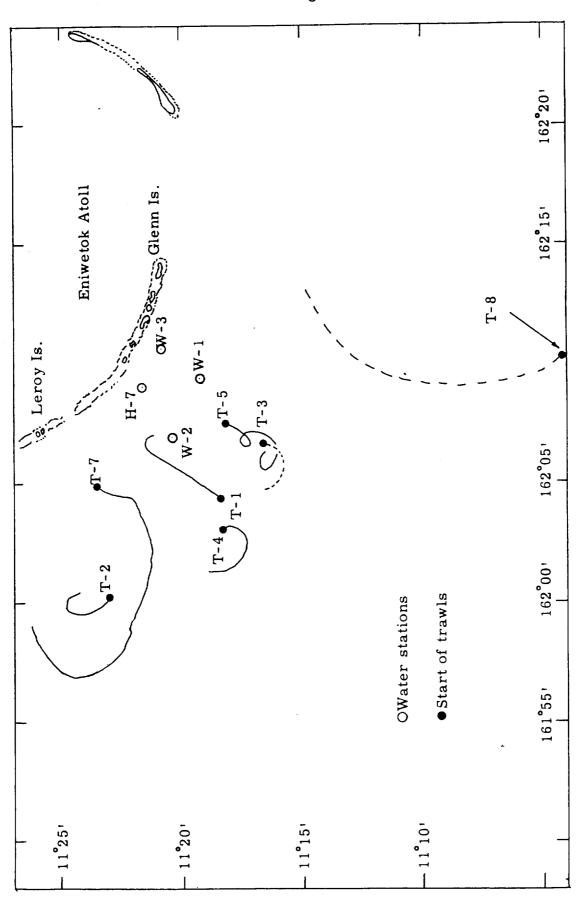


Figure 1. Area of the survey and location of the post-Wahoo plankton stations.



Pre-Wahoo plankton.(a) Location of plankton stations; (b) gross beta activity of plankton samples. Figure 2.



Location of the water stations and of the mid-water trawl hauls. The course plotted for T-8 is approximate. Figure 3.

bathythermograph (BT) winch, was lowered rapidly to the desired depth, towed at this depth for designated periods varying from 15 to 125 minutes, and then raised rapidly to deck level. Haul depths from the surface to 457 meters were determined by the use of a BT. The volume of the sample was determined volumetrically by allowing the sample to settle for five minutes in a graduate cylinder. Details of the plankton hauls are given in Appendix Table A.

In order to direct the movement of the ship while attempting to delineate the mass of radioactive water produced by the Wahoo detonation, a 5-cc portion of plankton samples P-22 to P-35 was filtered through a Millipore filter disc 47 mm in diameter (pore size 0.45μ), and the total gamma activity of the plankton was determined immediately on the wet sample. Later, at the Eniwetok Marine Biological Laboratory on Parry Island, the gross beta activity of the sample was determined on the dried material.

In some samples the plankton were separated into "large" and "small" plankton. The sample was first filtered through a bronze screen 2-3/4 inches in diameter with a mesh of 99 to the inch (large plankton), and then through a Millipore filter disc with a pore size of 0.45μ (small plankton).

For the determination of gross beta activity, a portion of each sample was placed on a 1-1/2-inch stainless steel planchet, dried under infrared lamps, and counted aboard ship. The beta-counting equipment consisted

of an end-window counter with a 2-inch Anton tube in a 3-inch Anton shield and a Nuclear-Chicago Model 181 scaler. The correction factors for geometry, self-absorption, backscatter, and decay used to convert c/m to d/m/g of wet tissue at time of collection are included in Appendix Tables D to H.

The samples for total gamma counting were placed in pliofilm bags and counted in a single-channel, 50-position, automatic-advance spectrometer with a 2-inch well-type sodium iodide crystal. Samples containing significant amounts of gamma activity were analyzed to determine the energies of the radionuclides present. Samples containing detectable amounts of beta activity were recounted at frequent intervals to determine radioactive decay.

Because of the difficulties of weighing samples aboard ship, the weights were determined at the Eniwetok Marine Biological Laboratory.

The radiochemical composition of nine samples was determined by ion-exchange techniques.

Fish

A mid-water trawl of the Isaacs-Kidd type with a spread of six

(1)
feet was used to collect fish and macroplankton from depths of 14
to 365 meters. Trawl hauls were made at speeds of 4 to 5 knots
and for periods of 35 to 147 minutes. A deep-sea, steam-powered
anchor-winch was used for streaming the trawl with a 1/2-inch

wire cable. The maximum tension obtained during the deepest trawl was approximately 6,000 pounds as measured by a dynamometer. The average tension while trawling was approximately 3,500 pounds. Details of each trawl haul are given in Appendix Table B.

After the small fish, shrimp, and miscellaneous plankton were separated, portions of each of these were placed on planchets for beta counting and in pliofilm bags for gamma counting. Radiochemical analyses were made of six samples by ion-exchange techniques.

Water

Four series of water samples, one before and three following the Wahoo detonation, were obtained from casts with Nansen bottles made by personnel of the U. S. Navy Hydrographic Office. The last three series consisted of seven samples obtained from depths of 0 to 300 meters, whereas the first series consisted of nineteen samples from depths of 0 to 1,039 meters. Details of each sample are given in Table 1.

Five hundred ml of water were passed through a Millipore filter 47 mm in diameter and with a pore size of 0.45 μ to collect the particulate matter present in the water. The filter paper was placed on a 1-1/2-inch stainless steel planchet, dried, and dissolved with acetone, then counted for total beta activity. Four ml of saturated sodium carbonate were added to 80 ml of the filtrate, and the carbonate precipitate was collected on a second Millipore filter paper and counted for beta activity. Naturally occurring K^{40} , normally present in sea water to the extent of

about 660 d/m/l (beta), remains in the filtrate by this method. Radiocesium also remains in the filtrate; however, it accounts for less than (3) one per cent of the total disintegration rate at the time of collection.

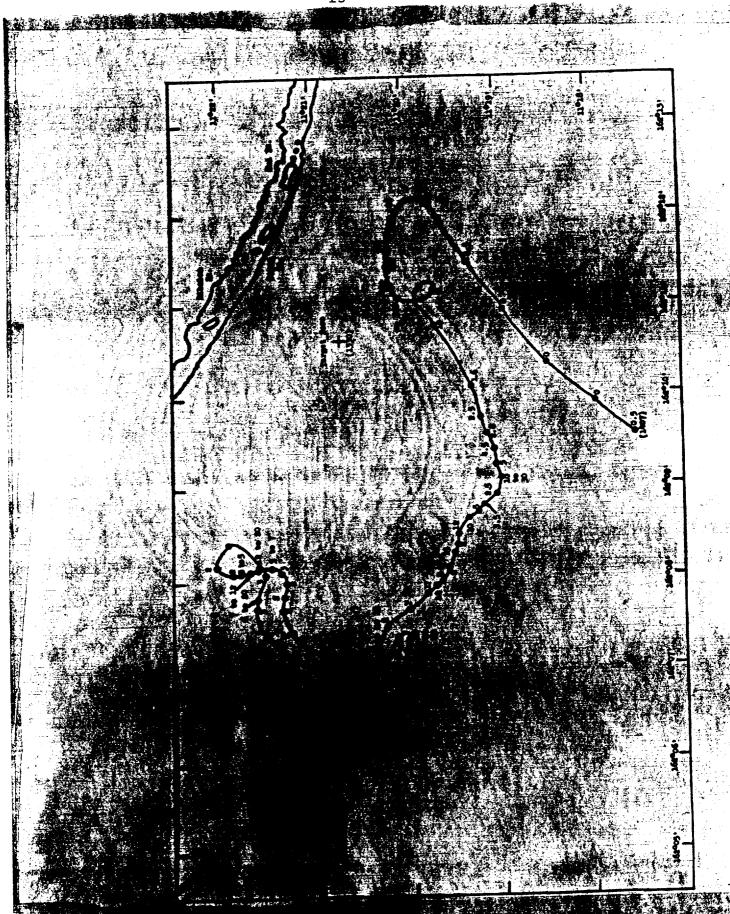
Gamma radiation survey of surface water

The gamma radiation measurements of the surface water immediately following the Wahoo test were obtained with U. S. Navy survey meters (AN/PDR-27C and AN/PDR-43(XN-2)) by three-man teams stationed on the bow of the ship. The readings were relayed to the bridge by telephone at frequent intervals and plotted with the positions of the ship during the survey. The ship was maneuvered to follow a 10 mr/hr surface radiation field (Fig. 4).

Radiochemical analyses of the samples

Radiochemical analyses by ion-exchange techniques were made of sixteen samples including plankton, fish, shrimp, and squid. The samples were wet-ashed with concentrated HNO3 and superoxol and dissolved in 0.2N HCL. The samples were then passed through 0.28 cm² x 10-cm columns containing Dowex-50* (100-200 mesh), and the radioisotopes were eluted with 50 ml of the following reagents: 0.5 per cent oxalic acid and 5 per cent ammonium citrate at pH 3, pH 4, and pH 6.5. (9) Aliquots of the collected fractions were counted for total gamma activity, and the fractions with significant amounts of gamma radioactivity were

^{*}Available from Dow Chemical Co., Midland, Michigan.



analyzed in the gamma spectrometer described above. The specific radioisotopes could be identified in a sample when the radioactivity of the isotope was one per cent or more of the total radioactivity of the sample. The fractions which contained more than one radioisotope were further separated by other ion-exchange methods.

Anion fractions

The anion fractions of plankton samples P-6, P-13 and P-15 + P-17 were dried, redissolved in 0.1 M oxalic acid, and passed through 0.13 cm² x 26-cm columns containing Dowex-1* (200-400 mesh) for the separation of the anions of Sb, Sn, Te, and Ru. The separation was done by the modified procedure of Smith and Reynolds as described by (13) Palumbo and Lowman.

The anion fractions of samples P-25 + P-26, P-24 + P-35, and
P-13 + P-15, and the fractions from the secondary separation of P-15 +
P-17, were dried and redissolved in concentrated HCl to separate the
(16)
complex radionuclide mixtures, as described by Wish. In these experiments the samples were passed through 0.03 cm² x 15 cm-columns
containing Dowex-1 (200-400 mesh) and the radionuclides were sequentially removed by using mixtures of hydrofluoric and hydrochloric
acids. Because of the corrosive nature of the HF acid, column plugs
made of lucite shavings were used and the fractions were collected in

^{*}Available from Dow Chemical Co., Midland, Michigan,

lusteroid test tubes.

Oxalate fractions

The oxalate fraction of the primary ion-exchange separation of sample P-6 was dried and redissolved in concentrated HCl for the separation of the transition elements, as described by Kraus and (6)

Moore. The sample was passed through a 0.28 cm² x 26-cm column containing Dowex-1-(200-400 mesh).

Determination of strontium-90

Samples of plankton and a sample of shrimp were analyzed for Sr⁹⁰ (5) by the ion-exchange method of Kawabata and Held. The ashed sample was first treated with 80 per cent HNO3 to remove strontium from the bulk of the sample and from most of the other radioisotopes present. The solution containing the strontium nitrate precipitate was filtered through a glass filter and was then dissolved in hot 0.2N HCl. After cooling, it was passed through a cationic resin column and possible contaminants were eluted from the column with 0.5 per cent oxalic acid and 5 per cent ammonium citrate at pH 3.5. The column was then stored for 14 days to allow the Y⁹⁰ daughter to build up. The Y⁹⁰ was reeluted from the column with ammonium citrate at pH 3.5 and the amount of Sr⁹⁰ present was calculated from the amount of Y⁹⁰ recovered. The identity of Y⁹⁰ was confirmed by determining the decay rate of the sample eluted.

RESULTS

Initial radiation survey

The Wahoo device was detonated at 1330 on May 16, 1958, and at 1407 hours, the radiation survey to determine the 10 mr/hr limits of the radioactive water mass was started. A summary of the ship's track and the radiation readings in mr/hr recorded during the survey are presented in Figure 4. Immediately before the Wahoo test while the ship was 7,000 yards south of target zero, the background radiation aboard ship was 2.0 mr/hr. The average background radiation in the target area was 1.5 mr/hr. In the following discussion the average background value has been subtracted from the values presented.

Following the detonation, as the ship proceeded in a northeasterly direction, it entered a 3 mr/hr field, at which time (1435) it was turned to the west, During the turn, the radiation level increased rapidly to 500 mr/hr (1451) and then dropped rapidly to 5 mr/hr as the ship's course was changed sharply southward and away from target zero. This point, approximately one mile southeast from target zero, probably represents the limit of the main body of radioactive water at this time (1455). The survey was continued in a southwesterly direction and at 1510, when readings of 10 to 30 mr/hr were obtained, the course was changed to a northwesterly one. At 1530, readings of 2 to 8 mr/hr were obtained

when the ship was three and one-quarter miles due west of target zero. As the ship continued northward and then circled back (1542) to parallel the course taken earlier, radiation readings of 2 to 20 mr/hr were observed. At the northernmost point of the survey the readings were 2 mr/hr, and at the westernmost point they were 1 mr/hr. This line represents the approximate detectable downwind limits of contamination at this time. At 1618 the initial survey was completed when the ship was approximately 6,500 yards downwind (WSW) from target zero.

The level of radioactivity in the target area during the next fortyeight-hour period decreased to the pre-Wahoo level.

Water

The gross beta radioactivity in the pre and post test samples is given in Table 1. The radioactivity in the samples collected prior to the Wahoo detonation was limited to the top 39 meters and for the most part was in a soluble form. Five and one-half hours after detonation, radioactivity was present at all depths sampled and the amounts of radioactivity were much higher than those found one and three-quarter miles southwest of target zero. The particulate matter suspended in the water accounted for 6 to 40 per cent of the gross beta activity. Twenty-seven hours after detonation, four miles due west of target zero, the measured amounts of beta radioactivity in the water were approximately the same. At W + 48, at target zero, the greatest

amount of radioactivity was found in the samples taken below the thermocline.

Plankton

The radioactivity measurements of the plankton samples are given A small amount of beta activity found in the first four in Table 2. samples can be attributed to residual contamination from previous tests. Because of the thousand-fold increase in radioactivity of plankton sample P-6 collected on May 6, 1958, as compared to those collected previous to this date, it is assumed that this increase in radioactivity was due to the fallout of the Cactus device detonated on the same day. Samples P-7 to P-12, collected before the Wahoo test, contained radioactivity in lower amounts (Fig. 2). These samples were examined by gamma spectrometry immediately after collection and the following short-lived isotopes were present in the decreasing order of their contribution to the $Mo^{99}-Tc^{99}m$ (66 hours),* Ce^{141} total activity: Np²³⁹ (2.3 days) Pr¹⁴¹ (33.1 days),* Ru ¹⁰³-Rh¹⁰³ (39.8 days),* Ba¹⁴⁰-La¹⁴⁰ (12.8 days)* $Te^{132}-I^{132}$ (77.7 hours)* $Zr^{95}-Nb^{95}$ (65 days)* $Ce^{144}-Pr^{144}$ (285 days), and Ru^{106} - Rh^{106} (1 year), in both the large and small plankton. A summary of the results of separations of the radionuclides in sample P-6 by ion-exchange methods is given in Table 3. Gamma spectrum analyses indicated that plankton samples P-7 to P-12 contained these radionuclides in the same relative proportions.

^{*}Effective half life.

Table 2. Radioactivity of the plankton. Values expressed as d/m/g of gross beta activity and c/m/cc of total gamma activity at time of collection.

			Radioactivity	
			Gross beta activity	Gamma activity
Sample	Collec	ted	10^6 d/m/g of	10^6 c/m/cc of
no.	Date	Hour	dry plankton*	wet plankton
P-1	5/4/58	1540	0.000374	
P-2	11	1945	0.000854	
P-3	5/5	1423	0.000346	
P-4	* ***	1950	0.000434	
P-6	5/6	1615	292.0	
P-7	11	1953	53.2	
P-8	5/7	1427	11.3 .	
P-9	#1	1637	4.06	
P-10	5/8	1845	8.58	
P-11	5/9	1225	0.424	
P-12	5/10	1300	0.00244	
P-13	5/16	1609	32.2	
P-14	11	19 07	257	
P-15	5/17	1015	4.76	
P-16	11	1405	10.3	
P-17	11	1705	96.2	
P-18	11	2140	17.4	
P-19	5/18	1040	25.2	
P-20	11	1830	15.5	
P-21	11	2304	33.7	
P-22	5/19	0858	18.6	0.200
P-23	11	1000	46.6	0.166
P-24	11	1100	3.60	0.182
P-25	11	1230		0.286
P-26	11	1352		0.220
P-27	*1	1814	7.66	0.100
P-28	†1	1935	12.1	0.125
P-29	* 1	2004	27.4	0.080
P-30	5/20	0101	7.52	0.0182
P-31	11	0145	6.93	0.0129
P-32	11	0400		0.0222
P-33	11	0617	3.18	0.0432
P-34	11	0855		0.0344
P-35	11	1510		0.0480
P-36	5/ 2 6	0945	2.30	

^{*}Correction factor for self-absorption based on K40.

Gamma emitting radioisotopes in samples of plankton. Values expressed as per cent of the total radioactivity contributed by each radioisotope at the time of collection. Table 3.

			ample			
		Percent	of tota	lacti	>	6
			P-13*	P-15	P-25	P-34
Radioisotope	P-6	P-13	P-15*	P-17	P-26	P-35
Np 239	56	73	29	62	81	74
Mo99-Tc99m	31	2.2	17	13	3.6	5.3
Ce141-Pr141	3,6	2.7	9.0	6.0	2.0	1.9
Ru103-Rh103	3.0	1.5	0.8	0.3	1.1	1.0
Ba140-La140	2.9	2.4	0.7	2.2	4.4	2.3
Te132-1132	1.4	7.3	13	12	5.4	6.6
Zr^{95} - Nb^{95}	1.1	pu	0.3	0.5	0.7	1.7
Ce144-Pr144	9.0	nd	pu	0.1	pu	pu
Ru106-Rh106	0.3	pu	pu	pu	pu	pu
U237	nd**	11	0.3	1.2	2.0	3,3
Ru ¹⁰⁵ -Rh ¹⁰⁵	pu	pu	pu	7.0	pu	pu
Total d/m/g x 10 ⁶	100	360	140	150	15.0	8.6

* Small plankton. ** Not detected.

On may 14, 1958, a second fallout was detected in the area in which the ship was operating. This contamination may have resulted from the Butternut, Fir, or Koa devices which were detonated one to to two days earlier. General contamination of the ship at a level of 5 to 10 mr/hr raised the background in the counting room to a very high level (445,000 c/m total gamma, 0.02-2 MEV). This level decreased to 60,000 c/m just prior to the Wahoo detonation on May 16, 1958.

The first post-Wahoo plankton sample, P-13, was taken at the end of the initial radiation survey (1609 hours) at a depth of 9 to 15 meters in an area with a gamma radiation level of 10 mr/hr, four miles west of target zero. The amount of radioactivity was high (32,000,000 d/m/g dry) but not as high as that found in sample P-14 taken that evening (1907 hours) at a position less than three miles to the southeast. These differences in amounts of radioactivity may reflect the variation in the radioactivity of samples taken in the target area or may indicate that maximum uptake of radioactivity by the plankton did not occur until this time. The location of the plankton sampling stations and the radioactivity of the samples collected after Wahoo are shown in Figures 1, 5, and 6.

The results of ion-exchange separations of the radionuclides of some of the plankton samples and analysis of the fractions by gamma spectrometry are given in Table 3. The percentages of total activity

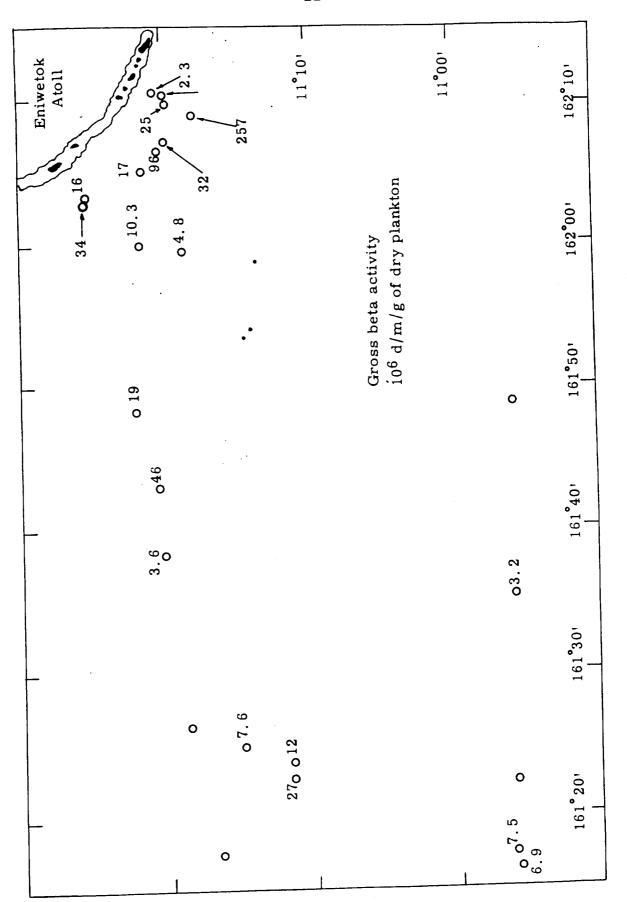


Figure 5. Gross beta activity of the post-Wahoo plankton samples.

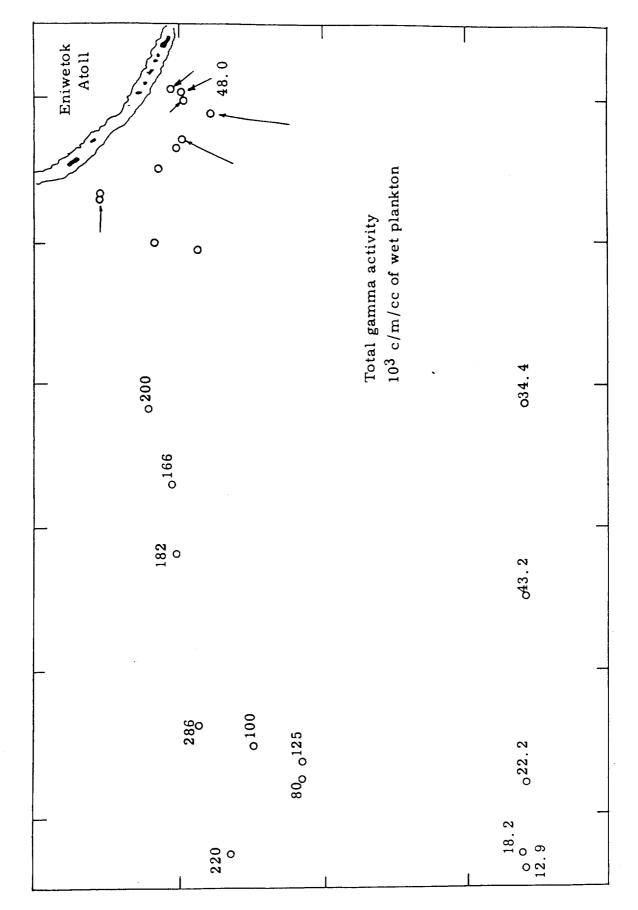


Figure 6. Total gamma activity of the post-Wahoo plankton samples.

given in Table 3 are based upon the calculated activity of the samples at the time of collection.

Levels of strontium-90

The results of the Sr⁹⁰ determinations of plankton and a shrimp sample are given in Table 4. The only pre-Wahoo plankton samples that contained detectable amounts of Sr⁹⁰ were samples P-6 and P-7, collected on the same day the Cactus device was detonated. Samples P-8 and P-12, collected 19 to 80 hours later, contained no Sr⁹⁰.

The plankton samples collected after Wahoo, however, contained levels of Sr^{90} that comprised from 0.0001 per cent to 0.04 per cent of the total beta radioactivity. Apparently the Wahoo detonation made a new source of Sr^{90} available to the plankton.

Fish and other organisms

Eight mid-water trawls were made during the survey, one of which was lost during the recovery operation. The following organisms were collected most frequently: euphausids, lantern fish, crab larvae, and miscellaneous macroplankton species. Eel larvae and small squid were obtained in two hauls. Silversides (fish of the family Atherinidae) and hatchet fish (Sternoptychidae) were obtained only once.

The gross beta radioactivity of the organisms collected in the midwater trawl is given in Table 5. There was no detectable radioactivity in the samples collected before Wahoo (T-1 to T-4). Post-Wahoo

Table 4. Sr⁹⁰ in samples collected aboard the USS Rehoboth May 6 to May 20, 1958. Analyses made January 21, 1959. Values expressed as d/m/g of dry weight at time of collection.

Sample			Date of collection	Gross beta activity 10 ⁶ d/m/g	Sr ⁹⁰ d/m/g	Sr ⁹⁰ per cent of gro beta activity
		1 1 . 4	E 6 E0	290	2050	0.00070
P-6	Large p	lankton	5-6-58		12	0.00002
P-7				55		
P-8	11	11	5-7	11	0	0
P-9	11	11	11	4.0	0	0
P-10	,11	11	5-8	8.6	0	0
P-12	11	11	5-10	0.0024	0	0
P-13	f 1	11	5-16	32	6.0	
P-13)	Small p	lankton	11	•		0.0000
P-15)	11	11	5-17	44.0	415	0.00090
P-15	Large p	lankton	5-17	4.8	1710	0.04000
P-16	11	11	11	10	441	0.00400
P-17	**	11	11	96	226	0.00020
P-18	11	11	ff	17	3280	0.02000
P-25)	13	11	5-19	54	705	0.00010
P-26)	11	11	11	34	103	0.00010
P-34)	ŧŧ	**	5-20	8.9	1030	0.01000
P-35)	11	11	11	•••		
T-7	Shrimp		5-18	3.2	0	0

Table 5. Gross beta radioactivity of the organisms collected in the mid-water trawls. Values expressed as d/m/g of dry weight at time of collection.

Sample no.	Maximum depth (meters)	Type of organism	d/m/g
T-1	180	Shrimp Plankton	Background** Background**
T-2	180	Miscellaneous*	Background**
T-3	175	Miscellaneous*	Background**
T-4	6	Miscellaneous*	Background**
T-5	25	Fish Shrimp Squid	259,000 2,680,000 3,060,000
T-7	365	Fish Shrimp Squid Plankton (large)	1,730,000 3,240,000 900,000 114,000
T-8	365	Fish Shrimp Plankton (large)	756,000 1,050,000 106,000

^{*}Mixture of small fish, shrimp and plankton.

^{**}Background, 16 c/m.

organisms, however, contained high amounts, and the radioactivity of the whole shrimp samples was approximately two to ten times as high as that of the whole fish.

Fish also were collected with three of the plankton samples, and two dolphins (Coryphaena hippurus) were caught by hook and line. The gross beta activity of these fish, of a squid, and of a portion of an alga (Turbinaria ornata*) collected during this period are given in Table 6.

Results of the radiochemical separations and gamma spectrum analyses of some samples in Table 6, as well as others, are given in Table 7.

Table 6. Gross beta radioactivity of the miscellaneous samples. Values expressed as d/m/g of dry weight at time of collection.

Sample No.	Organism	$d/m/g \times 10^3$
P-7	Lantern fish (whole)	4,940
P-17	Jack "	82.9
P-35	Blenny "	69.0
F-1	Dolphin	
	muscle	12.0
	liver	7.1
F-2	Squid	
· ·	muscle	62.6
•	stomach	156
A-1	Alga(Turbinaria ornata)	5,840

^{*}This alga is not pelagic and normally grows on the reef.

Table 7. Relative amounts of the radioisotopes present in fish, crustacea, squid, and in an alga collected May 6, 1958 to May 18, 1958.

Relative amount of radioisotope

		Te 13			Ba140				Ce ¹⁴¹
Organism	Np ²³⁹	1^{132}	Tc^{99m}	Rh ¹⁰³	La^{140}	Nb^{95}	Rh ¹⁰	6 _U 237	Pr ¹⁴¹
Fish*	++++	+++	+	++	+	+		+	
Shrimp*	++++	++		+++	+		+	++	
Squid*	++++	+++		+				+	
Fish**	++++	++	+	+	+			+	
Shrimp**	++++	+++							
Lantern fish**	++++	+++	+						
Jack	++++	+++	++					+	
Dolphin (liver)	++++	++		++					
Alga(Turbinaria									
ornata)					++				+++

^{+ =} low amount, 5 per cent of total activity; ++ = 6-20 per cent of total activity; +++ = 21-49 per cent of total activity; ++++ = 50 per cent of total activity.

^{*}Collected in mid-water trawl T-5; **collected in mid-water trawl T-7.

DISCUSSION

The extent of contamination of the water immediately following the Wahoo detonation was determined by means of gamma survey meters. This initial survey showed that at W + 1-1/2 (one and one-half hours after Wahoo) the area of contamination extended one and one-half miles southeast of the target, and at W + 2-1/2 it extended about four miles west of target zero. The radioactivity of water samples taken at W + 5, about two miles southwest of target zero, was found mostly above the thermocline, although some radioactivity was found in the deeper samples.

At W + 27, survey meter measurements taken four miles west of target zero were not significantly greater than the pre-Wahoo levels. However, gross beta activity determinations of water from the same area contained high levels of radioactivity in the uppermost 50 meters and lesser amounts in the deeper water. At W + 48, in water samples taken at target zero, the greatest amount of radioactivity was in the samples taken below the thermocline. These data show that the deeper water was moving horizontally at a slower rate than the water above the thermocline.

The radioactivity measurements of the plankton showed that in three days the contamination had extended at least 50 miles to the west of target zero. Samples taken 20 miles to the south, about 10 hours

later, contained the lowest levels of gamma radioactivity. The low levels in the plankton from this location indicated that the south-western perimeter of the contaminated water mass at this time was only a few miles away. The low level of radioactivity also suggests that the radioactive water had travelled primarily in a westward rather than a southwestward direction.

The radioisotopic composition of the plankton samples collected before and after the Wahoo detonation is given in Table 3. $\rm Np^{239}$ was the major contributor in all of the samples, but the relative amounts of $\rm Mo^{99}\text{-}Tc^{99m}$, $\rm Te^{132}\text{-}I^{132}$, and $\rm U^{237}$ differed, $\rm Mo^{99}\text{-}Tc^{99m}$ was much higher and $\rm Te^{132}\text{-}I^{132}$ was much lower in the pre-Wahoo sample. $\rm Ru^{105}\text{-}Rh^{105}$ was detected in one post-Wahoo sample and $\rm Ru^{106}\text{-}Rh^{106}$ was detected in one pre-Wahoo sample.

These differences in radioisotopic content may be used to establish the origin of the radioactive material in the samples collected during the survey. As seen in Table 3, the radioisotopes in samples P-25 and P-26, collected at the western periphery, were similar to the radioisotopes in earlier samples; therefore, it was assumed that the origin of the radioactive material was the same for all samples collected after the Wahoo detonation.

Results of radiochemical separations and gamma spectrum analyse revealed that the percentages of ${\rm Ba^{140}\text{-}La^{140}}$ and ${\rm U^{237}}$ were higher i

large plankton and ${\rm Mo}^{99}\text{-Tc}^{99{\rm m}}$ were higher in small plankton; otherwise the radioisotopic compositions were similar.

Analysis of sea water collected shortly after another underwater detonation (Umbrella), in the lagoon at Eniwetok, showed that Np 239 , Te 132 -I 132 , Ce 143 -Pr 143 , and Mo 99 -Tc 99m were the predominant radioisotopes in the particulate matter (>0.45 μ) present in the surface water near target zero (Table 8).

Table 8. Percentage of gamma emitting radioisotopes in the particulate ($> 0.45\mu$) and soluble ($< 0.45\mu$) form in sea water collected 75 minutes after the Umbrella event(8)

Radioisotope	Particulate	Per cent of particulate activity	Soluble 10 ⁶ d/m/l	Per cent of soluble activity
Np ²³⁹	5. 2	62.5	30.0	75.6
U ²³⁷	0.02	0.2	4.0	10.1
$Te^{132}-I^{132}$	1.9	22.3	1.9	4.7
Mo ⁹⁹ -Tc ^{99m}	0.48	5.8	3.1	7.8
Ce ^{l4l} -Pr ^{l4l}	0.028	0.3		0
$\mathrm{Ce}^{\mathrm{l}43}\text{-}\mathrm{Pr}^{\mathrm{l}43}$	0.66	7.9		0
Ce^{l44} - Pr^{l44}	0.0028	۷0.1		0
Ba ¹⁴⁰ -La ¹⁴⁰	0.076	0.9	0.7	1.8

Some of these isotopes also were present in high amounts in the $<0.45\mu$ fraction (Table 8), which could include materials

in the colloidal form. These radioisotopes, with the exception of Ce^{143} - Pr^{143} , were also the predominant ones in the plankton collected after Wahoo, a weapons testing device identical to Umbrella. These data suggest that in the first few days after detonation of a nuclear device these radioisotopes are mostly insoluble and that in this form they are removed from the water by the plankton.

In considering the distribution of radioactivity in the sea following a nuclear detonation, the role of the plankton should be considered. To estimate the relative amounts of radioactivity in water and plankton samples, the amount of water strained by the plankton net was calculated in the following manner.

The total volume of water filtered was calculated to be the product of the distance the net was towed, the filtering efficiency of the net (50 per cent), and its cross-sectional area. When the radioactivity of the total volume of water that passed through the net was calculated and compared with the radioactivity of the plankton in the same volume of water, the value for the water was found to be several thousand times greater (Table 9). This comparison indicates that the removal of radioactivity from the water by the plankton was apparently a minor factor, at least in the first 48 hours after detonation.

The amount of Sr⁹⁰ in the samples varied from 0 to 0.04 per cent of the gross beta activity. These values agree well with those obtained

Table 9. Comparison of the gross beta activity of the total plankton and total water strained for three samples collected after the Wahoo detonation.

			Total Water	volume Planktor	d/m/to n volur		d/m/total volume water
Sample	Date	Hour	M ³	cc	Water	Plankton	plankton
P-14	5/16/58	1907		8		1.7x10 ⁶	4, 400
W-1	5/16/58	1910	90		7.5x10 ⁹		
P-17	5/17/58	1705		14		5.2×10^6	11,000
W-2	5/17/58	1653	180		5.7x10 ¹⁰		
P-19	5/18/58	1040		10		1.1x10 ⁶	3,000
W-3	5/18/58	1348	180		3.3x10 ⁹		

in other investigations of the levels of Sr^{90} in marine organisms in the (3, 4, 7, 11) open sea .

The low levels of Sr^{90} in the plankton may be due to the low amounts present at the time of collection and to the unavailability of Sr^{90} , which may be explained on the basis of solubility, coprecipitation with calcium carbonate, competition with calcium, and isotopic dilution by stable (7) strontium.

The small fish collected after Wahoo had essentially the same radioisotopic composition as the plankton samples. A sample of dolphin liver obtained before Wahoo contained Np^{239} , Ru^{103} - Rh^{103} , and Te^{132} - I^{132} but no detectable amounts of the other radioisotopes which were found in the smaller fish and in the plankton.

Samples of whole shrimp and squid had high levels of Np 239 and relatively higher levels of Te 132 -I 132 than the plankton. However, they contained no detectable amounts of Mo 99 -Tc 99m .

Because these data indicate that some discrimination occurs in the uptake of radioisotopes by marine organisms, it would be of interest to determine which isotopes are incorporated or metabolized by these organisms. Some of the radioisotopes were concentrated rapidly by samples of mixed plankton species; knowledge of their concentration and fate in single species of plankton would be of interest.

SUMMARY

- 1. A survey of the radioactivity of sea water and marine organisms was conducted at the Eniwetok Test Site from May 3, 1958 to May 20, 1958, immediately before and after an underwater nuclear detonation.
- 2. Seventy-two hours after the underwater nuclear detonation, Wahoo, the radioactive water mass thus produced extended more than 50 miles westward from site zero, both above and below the thermocline.

- 3. The radioisotopic composition of the plankton and other marine organisms was determined by means of ion-exchange separations and gamma spectrum analyses.
- 4. Np²³⁹ was the major radioisotope present in all of the samples, contributing more than 50 per cent of the total radioactivity.
- 5. Other radioisotopes present in the plankton in the order of contribution were: Te^{132} - I^{132} , Mo^{99} - Tc^{99} m, Ba^{140} - La^{140} , U^{237} , Ce^{141} - Pr^{141} , Ru^{103} - Rh^{103} , Zr^{95} - Nb^{95} and Ce^{144} - Pr^{144} . Ru^{105} - Rh^{105} and Ru^{106} - Rh^{106} were detected only once.
- 6. Small whole fish contained essentially the same radioisotopes as the plankton. A liver sample from a large dolphin (fish) contained only Np^{239} , Ru^{103} - Rh^{103} , and Te^{132} - I^{132} .
- 7. Shrimp and squid samples concentrated Np²³⁹ and Te¹³²-I¹³² but contained no Mo⁹⁹-Tc^{99m}.

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APPENDIX

Appendix Table A. Log of the plankton samples collected aboard the USS Rehoboth during May 1958.

Depth of	thermocline (m)		100	110	06		85	80	110	100	105	125	105			*	85 *		*	105	80	*
. 연 면	surface		84	81.8	82.3		82.2	81.3	82	82.4	81.7	81.6	81.6	81.9		•	81.7			85	81.5	
	Haul depth		51	58	59.4		63	58	59.2	62	26	63	63					43		89	61.5	*
Depth of	plankton tow (m)		274	207	197	189	183	201	183	189	183	186	192	12 & 8	25	09	102	98	183	183	186	45 & 12
	သ		18	20	23	10	10	30	18	25	34	, 01	37	22	∞	20	15	14	7	10	22	15
į	Time (min)		35	95		45	18	43	36	38	29	48	125	37	23	30	63	55	15	35	60	36
	sition E. Long.		162 09.2	162 08.6	162 08.6	162 06.7	162 07.6	162 06.6	162 08.1	162 07.0	162 06.0	162 09.3	162 08.6	162 07.4	162 09.4	161 59.5	161 59.8	162 06.7	162 04.5	162 09.7	162 03.2	162 03.2
	Posit N. Lat.		11 20.4	11 16.1	11 21.0	11 21.6	11 23.1	11 19.9	11 22.6	11 24.4	11 24.2	11 21.4	11 21.3	11 20.0	11 19.3	11 18.7	11 21.4	11 20.5	11 21.4	11 20.0	11 25.8	11 25.6
•	Hour		1540	1945	1423	1950	1615	1953	1427	1637	1845	1225	1300	1609	1907	1015	1405	1705	2140	1040	1830	2304
;	Collected Date Hou	May	4	4	ည	ည	9	9	7	7	œ	6	10	16	16	17	17	17	17	18	18	18
	Sample no.		P-1	P-2	P-3	P-4	P-6	P-7	P-8	P-9	P-10	P-11	P-12	P-13	P-14	P-15	P-16	P-17	P-18	P-19	P-20	P-21

Appendix Table A. (continued)

Depth of	thermocline	(m).															80	
Temp. F	Haul at	depth surface		87*		85*	*98	****	.05	85.5*				u	85, 5*	82.1	53.4 81.7	
Depth of	plankton tow H	(m) de	30	30	30	46	49			30	3 & 6	46			49	52	, 253	
		၁၁	10	15	S	æ	12		10	10	18	15	12	4	25	6		
	Time	(mim)	15	15	15	15	15) L	15	18	15	15	17	18	18	20	36	
	tion	E. Long.	161 43.1		161 28.2	161 17.6	161 25 0		161 23.5	161 22.8	161 17.4	161 16.4	161 22.0	161 35.2	161 49.0	162 10.0	162 10.7	
	Position	N. Lat.	11 20.9	11 20.2	11 18.4	11 16.6	11 14 8	0.11	11 11.3	11 11.3	10 56.0	10 56.0	10 56.0	10 56.0	10 56.0	11 20.0	11 20.7	
	cted	Hour	1000	1100	1230	1352	1814	# 101	1935	2004	0101	0145	0400	0617	0855	1510	0945	
	Collected		19	19	19	19	10	61	19	19	20	20	20	20	20	20	26	
	Sample	no.	P-23	P-24	P-25	96-a	1 F	F-21	P-28	P-29	P-30	P-31	P-32	P-33	P-34	P-35	P-36	

*Faulty BT (bathythermograph), trace not clear.

Appendix Table B. Log of the mid-water trawl samples.

	Collected	ted				Maximum Cable	Cable	Tern	Terap. F	Depth of
Sample	2	Hour	Hour N. Lat.	E. Long.	Time (min)	depth (m)	length (m)	depth	Surface	(m)
T-1	j	1114	11 18.4	162 04.3	62	177	640	09	82	120
T-2	5/4/58	1252	11 23.0	162 00.2	86	177	640	59	82	06
T-3	5/4/58	2205	11 16.7	162 06.6	75	175	640	62	81.6	06
T-4	5/7/58	2240	11 18.3	162 03.1	35	9	66			
T-5	5/16/58	2215	11 18.2	162 07.5	65	24	146			
T-6	Lost					•				
L-1	5/18/58	1518	11 23.5	162 04.8	207	366	1830			
T-8	5/20/58 1306	1306	11 04.2	162 10.2	118	366	1830			

Appendix Table C. Log of the miscellaneous samples.

		;	•	Position	non	Depth
alume		Collected	eq	•	,	(= 1
ampro	Organism	Date	Hour	N. Lat.	N. Lat. E. Long.	(111)
7	Lantern fish	5/6/58	1953	11 19.9	162 06.9	201
	. Iack	5/17/58	1705	11 20.5	162 06.7	86
- L		5/20/58	1510	11 20	162 10	52
-35	Blenny	5/10/58	1640	11 11	162 37.6	surface
⊷. - Tr.	Dolphin	06 / 61 / 6			169 10	surface
با 1- 2	Squid	5/19/58	0200	11 20	104 10	
A - 1	Alga (Turbinaria	5/18/58	1800	11 26.7	162 00.3	suriace
	ornata)					

Appendix Table D. Radioactivity of the plankton. Correction factors used to convert counts per minute at time of initial count to d/m/g dry weight at time of collection.

					At	time of	collection	
Sample	In	Initial count	ıt	Decay		Dry		d/m/g***
no.	Date	Hour	c/m	factor	c/m	weight	c/m/g	dry weight
P-1	5/4		5	*	3	. 205	107	374
p-2	=		10	=	10	. 123	244	854
т. Д-3	5/5		14	=	14	. 141	66	346
P-4) -		16	=	16	. 129	124	434
D.6	5/11	0845	111, 416	*	1,500,000	.018	83, 300, 000	292,000,000
D-7	5/9	1355	ົດ	-	350,000	. 023	15,000,000	53, 200, 000
т Б-8	5/7	1605	`	Ξ	94,000	. 029	3, 240, 000	300,
о г	· } =	1950	· 6	=	33, 500	. 029	1, 160, 000	090
P-10	5/9	1350		Ξ	71,000	. 029	2, 450, 000	8, 580, 000
P-11	5/9	1720	4, 799	=	5, 200	. 043	121,000	
P-12	5/10	2155	26	Ξ	37	,053	869	2, 440
P-13	5/16	1715	84,602	=	92,000	.010	9, 200, 000	
P-14	=	2012	18, 749	=	22,000	. 0003		
P-15	5/17	1115	6,692	Ξ	6, 800	. 005	1, 360, 000	4, 760, 000
D. 16	5/17	1620	14, 508	Ξ	14,700	. 005	2, 940, 000	10, 300, 000
D-17	- -	1845	101, 598	=	110,000	. 004	27, 500, 000	96, 200, 000
P-18	=	2320	` ຕໍ	Ξ		. 005	4,980,000	17, 400, 000
P-19	Ξ	1940		=		. 016	7, 190, 000	25, 200, 000
P-20	Ξ	2047	57, 980	Ξ	62,000	. 014	4, 430, 000	15, 500, 000

*No decay factor used because of low counts and long half lives.

^{**}Decay extrapolated from decay curves.

^{***}Includes correction factor for self-absorption, geometry, and back scatter (3.5).

Appendix Table D. - continued

		1											
	d/m/g***	dry weight	33, 700, 000	18, 600, 000	46, 600, 000	3, 600, 000	7, 660, 000	12, 100, 000	27, 400, 000	7, 520, 000	6, 930, 000	3, 180, 000	2, 300, 000
collection		c/m/g	9, 620, 000	5, 300, 000	13, 300, 000	1, 030, 000	2, 190, 000	3, 460, 000	7,820,000	2, 150, 000	1, 980, 000	908, 000	657, 000
At time of collection	Dry	weight	800.	. 017	. 012	. 032	. 035	.034	.015	. 008	. 017	. 032	. 045
Ą		c/m	77,000	90,083	159, 365	330,242	76, 507	117, 493	117, 288	17, 194	33,653	29,052	29,549
	Decay	factor	*	Ξ	Ξ	=	Ξ	Ξ	=	=	Ξ	Ξ	Ξ
	ınt	c/m	64, 795	8, 341	14,756	30, 578	7,084	10,879	10,860	1,592	3, 116	2,690	3,888
	Initial count	Hour	0733	0220	0755	0800	0830	0840	0060	0910	0830	1100	1030
	H	Date	5/19	9/9	Ξ	Ξ	=	=	=	Ξ	Ξ	Ξ	5/30
	Sample	no.	P-21	P-22	P-23	P-24	P-27	P-28	P-29	P-30	P-31	P-33	P-36

^{**} Decay extrapolated from decay curves. *** Includes correction factor for self-absorption, geometry, and back scatter (3.5).

Appendix Table E. Radioactivity of the mid-water trawl organisms. Correction factors used to convert counts per minute at time of initial count to d/m/g dry weight at time of collection.

	***		000	000	000
	d/m/g***	0 0 0	2, 680, 000 2, 680, 000 3, 060, 000	1, 730, 000 3, 240, 000 900, 000 114, 000	756,000 1,050,000 106,000
At time of collection	c/m/g		74, 100 767, 000 873, 000	495,000 925,000 257,000 32,600	216,000 300,000 30,200
t time of	dry wt.		. 270 . 086 . 105	. 105 . 106 . 105	. 058 . 285 . 223
A	c/m		20,000 66,000 91,700	52, 000 98, 000 27, 000 4, 790	12, 550 (85, 500 6, 740
Decay	factor		1.28* 1.18* 1.005*	1.11* 1.15* 1.06* 1.15	7.5** 7.1** 7.1**
•	m/ 5	77	15, 678 65, 823 91, 271	46, 802 85, 536 25, 377 4, 166	1, 673 12, 038 949
	Initial count	Inou	0015	2215 2223 2220 1030	1638 " 1635
;	Init	Date	5/17	5/18	5/30
		Organism	Fish Shrimp	Squid Fish Shrimp Squid Macroplankton	Fish Shrimp
	Sample	no.	T-5	L-1	T-8

***Includes correction factor for self-absorption, geometry and back scatter (3.5). *Calculated from decay curves

to convert counts per minute at time of initial count to d/m/g dry weight at time of collection. Appendix Table F. Radioactivity of the miscellaneous organisms. Correction factors used

¥	d/m/g.r.	4, 940, 000 82, 900 69, 000	12, 000 7, 100	62, 600 156, 000	5, 840, 000
	q/1		20	000	
At time of collection	c/m/g	1, 410, 000 23, 700 19, 700	3, 440 2, 030	17, 900 44, 700	1, 670, 000
At time	dry wt.	. 049 . 098 . 150	. 192	. 179	. 027
	c/m	69, 000 2, 320 2, 950	086	3, 200 13, 000	45,000
Decay	factor*	62.0 1.03 3.97	4.55	1.01 1.06	1.00
	c/m	1, 112 2, 277 743	145 331	3, 155 12, 263	44, 708
Initial count	Hour	1645 1908 1641	1643 1300	0933 1046	1852
Init	Date	5/30/58 5/17/58 5/30	5/30 5/27	5/19	5/18
	Organism	Lantern fish Jack Blenny	Dolphin Muscle Liver	Squid Muscle Stomach	(Turbinaria ornata)
Sample	no.	P-7 P-17 P-35	표 1 - 1	F-2	A-1

* Calculated from decay curves.

^{**} Includes correction factor for self-absorption, geometry and backscatter (3.5).

Appendix Table G. Radioactivity of the water (particulate matter). Correction factors used to convert counts per minute at time of initial count to d/m/l at time of collection.

		d/m/1	63, 700	62,000	10, 200	22,000	3, 820		1,890	3, 100	30, 400	24 400			9, 900	7, 600	740	2, 500	3 460	0, 200	2, 340	3,040	ຕົ	64, 600	5, 280	000 06	•
collection	Volume	E	.350	. 500	=	=	=	4	.490	. 500	. 500	Ξ	Ξ	Ξ :	Ξ	=	=	Ξ	600		: :	=	Ξ	Ξ	=	=	
time of		d/m**	22, 300	31,000	່ ແລ	11 000	î -		924	1, 550	15, 200	19 910	12, 210	9,570	4,950	3,800	370	1,250	1 100	1, 130	1, 170	1, 520	1, 720	32, 300	Î o		14, 500
At		c/m	6, 760	9,410	1,550	2 220	,	280	280	469	4, 600	200	3, 100	2,900	1,500	1,150	112	380	u G	070	354	460	520	9 800	~		4, 400
	Decay	factor*	1.86	1 71		1 64	•	1.34	1.90	1.71	1 02	1 .	1.01	1.06	1.04	1.05		1.05		1.00	1,00	1,05				1.01	1.01
	nt	c/m	3 636	5,502		000	2, 030	431	147	274	4 400		3, 698	2, 743			108	361	!	525	354	437	497	0010	L r	167	4,320
	tial count	Hour	9317		0.00	0167		0005	=	0200	1695	1035	1630	1757	1820					1600	1520	Ξ	1540	000	0001	1500	Ξ
	Initia	Date	5/16	-	=	. :		5/17	Ξ	Ξ	, n) I / C	=	Ξ	Ξ	Ξ	5/24	5/17		5/18	Ξ	Ξ	=	Ξ	: :	Ξ	=
	Depth	(m)		ט ע	n (25	20	100	200	300		O	5 8	48	7.9	3 -0	145	293		0	30	ָ ה ה	1 C	- ·	100	150	300
	Sample	no.	-	V-1-W	4TC ::	515	516	" 517	11 518	" 519		W - 2 - 520	521	11 522	1 7 9 2	12 12 12 13 13	- - - - - -	1, 526		W-3-527	1, 528	1 10 10 10 10 10	640		531	532	11 533

**Includes correction factor for self-absorption, geometry and backscatter (3.3). *Calculated from decay curves

Appendix Table H. Radioactivity of the water (filterable portion less K^{40}). Correction factors used to convert counts per minute at time of initial count to d/m/l at time of collection.

		d/m/l	156,000	16,	73 200	000	29, 900	22, 500	30, 500		000	165,000	152,000	141,000	60 000	000 66	23, 900	26,000	32,000			11,800	14, 100	11,000	10,900	73, 500	12,900	000 00		
collection	Volume	(1)	100	100		: :	=	Ξ	=			080.	=		Ξ	:	=	Ξ	Ξ			080.	Ξ	=	=	Ξ	=	: =	:	
At time of		4/m/p	15,600	ĵ -	-	7, 320	2, 990	2, 250	3,050			13,200	12, 200	î -	.	4, 800	1,910	2,080	2, 560	•		944	1, 130		872	2 0	5, 880	1, 030	3,920	
ď	i	æ/ 0	000	3, 900	2, 900	1,830	748	18.00 18.00 18.00	762] ;		3, 300	2, 040	3,040	2, 830	1, 200	. 477	520	640			936	0000	000	0770	212	1,470	258	981	
		Decay	Tactor	1.42	2.02	1,72	1 79	1 - 1	1. (2	1.12	· ·	1 05	1.00	•	1.18	1 02	•	1.10	1.00			Ċ		1.03			1.02	,	$\frac{1}{1,10}$	
		ı	-1	2, 757	1,438	1,065	1,000	435	327	443		101	5, 155	2, 578	2, 402	1 180	1, 100	404	337	288		(180	273	200	198	1 438	•	892))
		Initial count	Hour	2332	=		6700	=	0200	0210	0200	6	1828	1824	1919		0081	1915	1922	1913			1630	1650	1705	1650	1645	0 6 9 7	1645	2 1 1 1 1 1 1 1 1 1 1
		Ini	Date	5/16	=	1	5/17	=	=	=	=		5/17	Ξ	=	=	Ξ	=	Ξ	Ξ			Ξ	=	Ξ	=	=	. =	: =	
		Depth	(H)	c) (ဂ	25	50	100	200	300		0	90	67	48	7.2	96	145	293)) 		0	30) K	י כ	C).	100	150	300
		Samule	or direct	٠ از	C1C-I-M	1 514	" 515	" 516	" 517	" 518	" 519		W-2-520	010 M		" 522	'' 523	524	11 525	11 526	3		W-3-527	: 10 E	040			11 531		11 533

**Includes correction factor for self-absorption, geometry and backscatter (4.0). *Calculated from decay curves.